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Amendments to the Claims

This listing of claims will replace all prior versions, and listings of claims in the application:

Listing of Claims:

Claims 1-2 (Cancelled).

3(Currently amended). A transgenic mouse having integrated in its genome a nucleic acid construct according to elaim 1, comprising a mammalian T-cell lineage specific, expression regulatory sequence promoter operably linked to a mammalian Glucocorticoid Induced Leucine-Zipper (GILZ) GILR cDNA sequence, wherein said mouse expresses GILR GILZ in its T-cell lineage at an elevated level compared to a non-transgenic mouse and wherein the expression of GILR results in an alteration of the thymocyte subset composition and of caspase 3 activation.

4 (Currently amended). The transgenic mouse according to claim 3, wherein said mammalian T-cell lineage specific, expression regulatory sequence promoter comprises a human CD2 promoter and a human CD2 locus control region.

Claims 5-16 (Cancelled).

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17(Original). A method for screening compounds having glucocorticoid-related effects, comprising:

administering a potential candidate compound to a transgenic mouse of claim 3, and to a control non-transgenic mouse; and

determining whether said potential candidate compound exhibits glucocorticoid-related effects by comparing the effects of the administration of said potential candidate to said transgenic mouse and to said control non-transgenic mouse.

18 (Original). A method for screening compounds having glucocorticoid-related effects, comprising:

administering a potential candidate compound to a transgenic mouse of claim 4, and to a control non-transgenic mouse; and

determining whether said potential candidate compound exhibits glucocorticoid-related effects by comparing the effects of the administration of said potential candidate to said transgenic mouse and to said control non-transgenic mouse.

19 (Currently amended). A method of producing a transgenic mouse whose genome comprises a nucleic acid construct, wherein said construct comprises a mammalian T-cell lineage specific, expression regulatory sequence promoter operably linked

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to a <u>mammalian Glucocorticoid Induced leucine-Zipper (GILZ) GILR</u> cDNA sequence, said method comprising:

transferring a nucleic acid construct according to claim 1, comprising a mammalian T-cell lineage specific promoter operably linked to a GILR mammalian GILZ cDNA sequence to a fertilized mouse oocyte;

allowing the zygote resulting from the fertilized mouse oocyte to develop to term, thereby obtaining a transgenic mouse whose genome comprises the nucleic acid construct;

breeding said transgenic mouse with a non-transgenic mouse to generate offspring; and

selecting from the offspring a transgenic mouse whose genome comprises the nucleic acid construct, wherein said transgenic mouse expresses GILR in the T-cell lineage at an elevated level compared to a non-transgenic mouse.

Claim 20 (Cancelled).